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(71) Applicant (for all designated States except US):
MERCK PATENT GMBH [DE/DE]; Frankfurter Strasse 250,
D-64293 Darmstadt (DE).

(72) Inventors; and

(75) Inventors/Applicants (US only): JONCZYK, Alfred
[DE/DE]; Scheppallee 57, D-64295 Darmstadt (DE).
DIEFENBACH, Beate [DE/DE]; Curd-Jürgens-Strasse 2,
D-81739 Munich (DE). GOODMAN, Simon [GB/DE];
Friedrich-Ebert Strasse 102 A, D-64347 Griesheim (DE).
GROTH, Ulrich [DE/DE]; Jakobstrasse 39, D-78464
Konstanz (DE). ZISCHINSKY, Gunther [AT/DE];
Mangoldstrasse 26, D-78462 Konstanz (DE).

(74) Representative: MERCK PATENT GMBH; D-64271
Darmstadt (DE).

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(54) Title: CYCLIC PEPTIDE DERIVATIVES AS INHIBITORS OF INTEGRIN $\alpha_v\beta_6$

(54) Bezeichnung: CYCLISCHE PEPTIDDERIVATE ALS INHIBITOREN DES INTEGRINS $\alpha_v\beta_6$

(57) Abstract: The invention relates to novel peptide derivatives of formula (I): Cyclo-(Arg-X¹-Asp-X²-X³-X⁴-X⁵-X⁶-R¹); which are biologically active as ligands of integrin $\alpha_v\beta_6$, X¹ representing Ser, Gly or Thr; X² representing Leu, Ile, Nle, Val or Phe; X³ representing Asp, Glu, Lys or Phe; X⁴ representing Gly, Ala or Ser; X⁵ representing Leu, Ile, Nle, Val or Phe; X⁶ representing Arg, Har or Lys; and R¹ being left out or representing one or more ω -aminocarboxylic acid radicals, said ω -aminocarboxylic acid radical(s) being 500 to 2500 pm in length. Said amino acids can also be derivatised and the D and L forms of the optically active amino acid radicals are enclosed. The invention also relates to the physiologically suitable salts and solvates of the inventive derivatives.

(57) Zusammenfassung: Die Erfindung beschreibt neuartige Peptidderivate der Formel (I), welche als Liganden des Integrins $\alpha_v\beta_6$ biologisch wirksam sind: Cyclo-(Arg-X¹-Asp-X²-X³-X⁴-X⁵-X⁶-R¹), worin X¹ Ser, Gly oder Thr, X² Leu, Ile, Nle, Val oder Phe, X³ Asp, Glu, Lys oder Phe, X⁴ Gly, Ala oder Ser, X⁵ Leu, Ile, Nle, Val oder Phe, X⁶ Arg, Har oder Lys, R¹ fehlt oder einen oder mehrere ω -Aminocarbonsäurerest(e), wobei der oder die ω -Aminocarbonsäurerest(e) eine Länge von 500 bis 2500 pm aufweisen, bedeuten, wobei die genannten Aminosäuren auch derivatisiert sein können, die D- als auch die L-Formen der optisch aktiven Aminosäurereste eingeschlossen sind, sowie deren physiologisch unbedenklichen Salze und Solvate.

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